

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant: Ken LIPSON et al.

Title: METHODS OF MODULATING C-KIT TYROSINE PROTEIN KINASE FUNCTION WITH INDOLINONE COMPOUNDS

Prior Appl. No.: 09/741,842

Prior Appl. Filing Date: 12/22/2000

Examiner: Unassigned

Art Unit: Unassigned

INFORMATION DISCLOSURE STATEMENT
UNDER 37 CFR §1.56

Mail Stop PATENT APPLICATION
Commissioner for Patents
PO Box 1450
Alexandria, Virginia 22313-1450

Sir:

Applicants submit herewith on Form PTO/SB/08 a listing of the documents cited by or submitted to the U.S. PTO in parent application Serial No. 09/741,842, filed 12/22/2000. As provided in 37 CFR §1.98(d), copies of the documents are not being provided since they were previously submitted to the United States Patent & Trademark Office in the above-identified parent application.

The submission of any document herewith, which is not a statutory bar, is not intended as an admission that such document constitutes prior art against the claims of the present application or that such document is considered material to patentability as defined in 37 CFR §1.56(b). Applicants do not waive any rights to take any action which would be appropriate to antedate or otherwise remove as a competent reference any document which is determined to be a *prima facie* art reference against the claims of the present application.

TIMING OF THE DISCLOSURE

The listed documents are being submitted in compliance with 37 CFR §1.97(b), within three (3) months of the filing date of the application.

RELEVANCE OF EACH DOCUMENT

All of the documents are in English.

Applicants respectfully request that the listed documents be considered by the Examiner and be made of record in the present application and that an initialed copy of Form PTO/SB/08 be returned in accordance with MPEP §609.

The Commissioner is hereby authorized to charge any additional fees which may be required regarding this application under 37 CFR §§ 1.16-1.17, or credit any overpayment, to Deposit Account No. 19-0741. Should no proper payment be enclosed herewith, as by a check being in the wrong amount, unsigned, post-dated, otherwise improper or informal or even entirely missing, the Commissioner is authorized to charge the unpaid amount to Deposit Account No. 19-0741.

Date 6/23/03

FOLEY & LARDNER
Customer Number: 22428



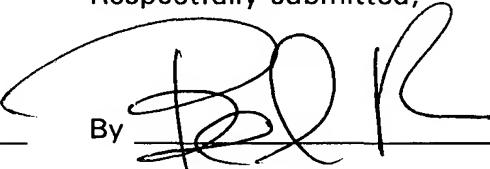
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PATENT TRADEMARK OFFICE

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Respectfully submitted,

By


Beth A. Burrous
Attorney for Applicant
Registration No. 35,087

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Group Art Unit	Unassigned
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Attorney Docket Number

038602-1607

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
		Number	Kind Code ² (if known)			
	A1	2,622,980		Copeland	12-23-1952	
	A2	2,872,372		Hull	02-03-1959	
	A3	2,968,557		Burgandt et al.	01-17-1961	
	A4	3,140,180		Fritz	07-07-1964	
	A5	3,308,134		Plostneiks	03-07-1967	
	A6	3,551,571		Pachter et al.	12-29-1970	
	A7	3,564,016		Schoen et al.	02-16-1971	
	A8	3,715,364		Hoff	02-06-1973	
	A9	3,880,871		Haugwitz et al.	04-29-1975	
	A10	3,922,163		Church et al.	11-25-1975	
	A11	4,002,643		Carson	01-11-1977	
	A12	4,002,749		Rovnyak	01-11-1977	
	A13	4,053,613		Rovnyak et al.	10-11-1977	
	A14	4,070,366		Gregorovich et al.	01-24-1978	
	A15	4,259,345		Cross et al.	03-31-1981	
	A16	4,259,346		Helmut STÄHLE et al.	03-31-1981	
	A17	4,343,923		Lenox et al.	08-10-1982	
	A18	4,376,110		David et al.	03-08-1983	

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		Number	Kind Code ² (if known)			
	A19	4,436,892		Zondler et al.	03-13-1984	
	A20	4,489,089		Wright, Jr. et al.	12-18-1984	
	A21	4,493,642		Furazawa et al.	01-15-1985	
	A22	4,493,842		Kunihiko FURUZAWA et al.	01-15-1985	
	A23	4,628,105		Schmid et al.	12-09-1986	
	A24	4,642,309		Michel et al.	02-10-1987	
	A25	4,826,847		Michel et al.	05-02-1989	
	A26	4,853,403		Shiraishi et al.	08-01-1989	
	A27	4,853,404		Takamura et al.	08-01-1989	
	A28	4,868,304		Larock	09-19-1989	
	A29	4,924,000		Rentzea et al.	05-08-1990	
	A30	4,966,849		Vallee et al.	10-30-1990	
	A31	4,971,996		Shiraishi et al.	11-20-1990	
	A32	4,987,146		Rohde et al.	01-22-1991	
	A33	5,043,348		Zoller et al.	08-27-1991	
	A34	5,043,454		Wriede et al.	08-27-1991	
	A35	5,047,554		Ehrgott et al.	09-10-1991	
	A36	5,051,417		Nadler et al.	09-24-1991	

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		Number	Kind Code ² (if known)			
	A37	5,057,538		Shiraishi et al.	10-15-1991	
	A38	5,082,856	A	Taniguchi et al.	01-21-1992	
	A39	5,082,856	A	Masao TANIGUCHI et al.	01-21-1992	
	A40	5,089,516	A	Shiraishi et al.	02-18-1992	
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	A42	5,145,983	A	West	09-08-1992	
	A43	5,153,217	A	Taniguchi et al.	10-06-1992	
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	A45	5,202,341	A	Shiraishi et al.	04-13-1993	
	A46	5,206,261	A	Kawaguchi et al.	04-27-1993	
	A47	5,217,999	A	Levitzki et al.	06-08-1993	
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	A49	5,278,184	A	Artico et al.	01-11-1994	
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	A52	5,322,950	A	Sircar et al.	06-21-1994	
	A53	5,330,992	A	Eissenstat et al.	07-19-1994	
	A54	5,332,736	A	Carmosin et al.	07-26-1994	

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		Number	Kind Code ² (if known)			
	A55	5,374,652	A	Buzzetti et al.	12-20-1994	
	A56	5,382,593	A	Le Baut et al.	01-17-1995	
	A57	5,389,661	A	Sircar et al.	02-14-1995	
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	A59	5,409,930	A	Spada et al.	04-25-1995	
	A60	5,409,949	A	Buzzetti et al.	04-25-1995	
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	A62	Re. 35,096	E	Taniguchi et al.	11-21-1995	
	A63	5,565,324	A	Still et al.	10-15-1996	
	A64	5,610,173	A	Schwartz et al.	03-11-1997	
	A65	5,723,665	A	Kato et al.	03-03-1998	
	A66	5,786,488	A	Tang et al.	07-28-1998	
	A67	5,792,783		tang et al.	08-11-1998	
	A68	5,792,783	A	Tang et al.	08-11-1998	
	A69	5,834,504	A	Tang et al.	11-10-1998	
	A70	5,849,710	A	Battistini et al.	12-15-1998	
	A71	5,880,141		tang et al.	03-09-1999	
	A72	5,880,141	A	Tang et al.	03-09-1999	

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		Number	Kind Code ² (if known)			
	A73	5,883,113	A	Tang et al.	03-16-1999	
	A74	5,883,116	A	Tang et al.	03-16-1999	
	A75	5,886,020	A	Tang et al.	03-23-1999	
	A76	Re. 36,256	E	Spada et al.	07-20-1999	
	A77	6,130,239	A	Chen et al.	10-10-2000	
	A78	6,133,305	A	Tang et al.	10-17-2000	
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	A81	6,395,736	B1	Thomas PARKS et al.	05-28-2002	
	A82	6,451,838	B1	Malcolm Wilson MOON et al.	09-17-2002	
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		Office ³	Number ⁴	Kind Code ⁵ (if known)				
	A84	WO	88/07035	A1	KANEYAFUCHI KAGAKU KOGYO KABUSHIKI KAISHA	09-22-1988		
	A85	WO	91/13055	A2	FARMITALIA CARLO ERBA SRL	09-05-1991		
	A86	WO	91/15495	A1	PFIZER INC.	10-17-1991		

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	A87	WO	92/03736	A1	SEIKAGAKU KOGYO KABUSHIKI KAISHA	03-05-1992		
	A88	WO	92/07830	A2	PFIZER INC.	05-14-1992		
	A89	WO	92/20642	A1	RHONEPOULENC RORER INTERNATIONAL	11-26-1992		
	A90	WO	92/21660	A1	PFIZER INC.	12-10-1992		
	A91	WO	93/01182	A1	FARMITALIA CARLO ERA SRL	01-21-1993		
	A92	WO	93/23040	A1	MERCK & CO., INC.	11-25-1993		
	A93	WO	94/03427	A1	WARNER-LAMBERT COMPANY	02-17-1994		
	A94	WO	94/10202	A1	GENENTECH, INC.	05-11-1994		
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	A102	WO	96/22976	A1	PHARMACIA S.P.A.	08-01-1996		
	A103	WO	96/32380	A1	PHARMACIA S.P.A.	10-17-1996		

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	A104	WO	96/40116	A1	SUGEN, INC.	12-19-1996	
	A105	WO	97/25986	A1	TAIHO PHARMACEUTICAL CO., LTD.	07-24-1997	
	A106	WO	97/34920	A1	SUGEN, INC.	09-25-1997	
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	A119	WO	99/61422	A1	SUGEN, INC.	12-02-1999	
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	A121	WO	00/08202	A2	SUGEN, INC.	02-17-2000	

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Date Submitted: June 23, 2003

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Prior Application Number	09/741,842
Prior Appl. Filing Date	12/22/2000
First Named Inventor	Ken Lipson
Group Art Unit	Unassigned
Examiner Name	Unassigned

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Attorney Docket Number 038602-1607

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	Foreign Patent Document			Name of Patentee or Applicant of Cited Documents	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T ⁶
		Office ³	Number ⁴	Kind Code ⁵ (if known)				
	A122	WO	00/38519	A1	SUGEN, INC.	07-06-2000		
	A123	WO	00/56709	A1	SUGEN, INC.	09-28-2000		
	A124	WO	01/60814	A2	SUGEN, INC.	08-23-2001		
	A125	DE	878,539		Von FREYBERG, et al.	06-05-1953		
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	A132	EP	0 252 713	B1	PFIZER INC.	01-13-1988		
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	A134	EP	0 351 213	A2	LES LABORATOIRES BEECHAM S.A.	01-17-1990		
	A135	EP	0 525 472	A2	FARMITALIA CARLO ERBA SRL	02-03-1993		
	A136	EP	0 566 226	B1	ZENECA LIMITED	10-20-1993		
	A137	EP	0 580 502	B1	ADIR ET COMPAGNIE	01-26-1994		
	A138	EP	0 626 377	B1	SHIONOGI & CO., LTD.	11-30-1994		

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				First Named Inventor	Ken Lipson
				Group Art Unit	Unassigned
				Examiner Name	Unassigned
Sheet	9	of	31	Attorney Docket Number	038602-1607

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		Office ³	Number ⁴	Kind Code ⁵ (if known)				
	A139	EP	0 632 102	A1	BAYER AG	01-04-1995		
	A140	EP	0 662 473	A1	PHARMACIA S.P.A.	07-12-1995		
	A141	EP	0 788 890	A1	AGFA-GEVAERT	08-13-1997		
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First Named Inventor	Ken Lipson
Group Art Unit	Unassigned
Examiner Name	Unassigned

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		Office ³	Number ⁴	Kind Code ⁵ (if known)				
	A156	AU	286870		IMPERIAL CHEMICAL INDUSTRIES OF AUSTRALIA AND NEW ZEALAND LIMITED	05-11-1967		

NON PATENT LITERATURE DOCUMENTS

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	A157	ABRAMOVITCH and HEY, "Internuclear cyclisation. Part VIII. Naphth[3:2:1-cd]oxindoles," <u>J. Chem. Soc.</u> , 1697-1703 (1954), Strand, London	
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	A159	Beilstein Reg. No. 236050, Beilstein Reference No. 4-21-00-06355	
	A160	AKBASAK and SUNAR-AKBASAK, "Oncogenes: cause or consequence in the development of glial tumors," <u>J. Neurol. Sci.</u> 111:119-133 (1992) © Elsevier Science Publishers	
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	A164	ANDREANI et al., "Synthesis and cardiotonic activity of 2-indolinones," <u>Chemical Abstracts</u> , Vol. 113, abstract no. 78106 (1990)	
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Prior Application Number	09/741,842
Prior Appl. Filing Date	12/22/2000
First Named Inventor	Ken Lipson
Group Art Unit	Unassigned
Examiner Name	Unassigned

Attorney Docket Number 038602-1607

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Group Art Unit	Unassigned
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	A185	BONNER et al., "Structure and Biological Activity of Human Homologs of the <i>raf/mil</i> Oncogene," <u>Molecular and Cellular Biology</u> 5:1400-1407 (1985) © The American Society for Microbiology	
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(use as many sheets as necessary)				First Named Inventor	Ken Lipson
Sheet	15	of	31	Group Art Unit	Unassigned
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	A208	ELLIOTT and RIVERS, "Reduction of Some Oxindolylidene Derivatives to 3-Substituted Oxindoles by Sodium Borohydride," <u>J. Med. Chem.</u> 29:2438-2440 (1964)			
	A209	ELLIOTT et al., "1-methyl-2-(3-oxindolenmethyl)-pyridinium," <u>Journal of Organic Chemistry</u> 29:2438-2440 (1964) DATABASE CROSSFIRE, Beilstein Reference No. 5-24			
	A210	FANTL et al., "Distinct Phosphotyrosines on a Growth Factor Receptor Bind to Specific Molecules That Mediate Different Signaling Pathways," <u>Cell</u> 69:413-423 (1992) © Cell Press			

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Prior Application Number	09/741,842
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	A215	FLOEGE et al., "Heparin suppresses mesangial cell proliferation and matrix expansion in experimental mesangioproliferative glomerulonephritis," <u>Kidney International</u> 43:369-380 (1993) © International Society of Nephrology
	A216	FOLKMAN and SHING, "Angiogenesis," <u>J. Biol. Chem.</u> 267:10931-10934 (1992) © The American Society for Biochemistry and Molecular Biology
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	A218	FOLKMAN, "Tumor Angiogenesis: Therapeutic Implications," <u>New England J. Medicine</u> 285:1182-1186 (1971)
	A219	FOLKMAN, "What is the Evidence that Tumors are Angiogenesis Dependent?" <u>Journal of the National Cancer Institute</u> 82:4-6 (1990)

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	A220	FOLKMAN, "Angiogenesis in Psoriasis: Therapeutic Implications," <u>J. Invest. Dermatol.</u> 59:40-43 (1973) copyright The Williams & Wilkins Co.		
	A221	GAZIT et al., "Tyrphostins. 2. Heterocyclic and α -Substituted Benzylidenemalononitrile Tyrphostins as Potent Inhibitors of EGF Receptor and ErbB2/neu Tyrosine Kinases," <u>J. Med. Chem.</u> 34:1896-1907 (1991) copyright Am. Clem. Soc.		
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	A223	GOLDRING and GOLDRING, "Cytokines and Cell Growth Control," <u>Critical Reviews in Eukaryotic Gene Expression</u> 1:301-326 (1991)		
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	A227	HEWGILL and STEWART, "Phenanthrene-4,5-quinones: a Synthesis of Morphenol," <u>J. Chem. Soc. Perkin Trans. I</u> :1305-1311 (1988)		
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	A265	MARTIN-LEON et al., "On the Cyclization to the Elusive Amino-4H-pyran Ring Some New Facts," <u>Liebigs Ann. Chem.</u> 101-104 (1990) copyright VCH Veilaks of Sellschaft mbH ©VCH		
	A266	MEL'NIKOVA TV et al., "Indole chemistry. XXXVIII. Cleavage of a carbon-carbon bond during the reaction of 2-aminoindoles with difunctional compounds," <u>Chemical Abstracts</u> 80 (1974) Abstract No. 003413		
	A267	MILLAUER et al., "High Affinity VEGF Binding and Developmental Expression Suggest Flk-1 as a Major Regulator of Vasculogenesis and Angiogenesis," <u>Cell</u> 72:835-846 (1993) © Cell Press		
	A268	MOHAMMADI et al., "Structures of the Tyrosine Kinase Domain of Fibroblast Growth Factor Receptor in Complex with Inhibitors," <u>Science</u> 276:955-960 (1997) © American Association for the Advancement of Science		
	A269	MORETO et al., "Study of the Laxative Properties of the Disodium Salt of the Sulfuric Diester of 3,3 Bis-(4-Hydroxyphenyl)-7-Methyl-2-Indolinone (DAN-603) in the Rat," <u>European Journal of Pharmacology</u> 36:221-226 (1976) ©North-Holland Publishing Company		
	A270	MORETO et al., "3,3-Bis-(4-Hydroxyphenyl)-7-Methyl-2-Indolinone (BHMI), the Active Metabolite of the Laxative Sulisatin," <u>Arzneimittel-Forschung Drug Research</u> 29:1561-1564 (1979)		
	A271	MORRISON et al., "Signal Transduction From Membrane to Cytoplasm: Growth Factors and Membrane-Bound Oncogene Products Increase Raf-1 Phosphorylation and Associated Protein Kinase Activity," <u>Proc. Natl. Acad. Sci. USA</u> 85:8855-8859 (1988)		
	A272	MOSMANN, "Rapid Colorimetric Assay for Cellular Growth and Survival: Application to Proliferation and Cytotoxicity Assays," <u>J. Immunol. Methods</u> 65:55-63 (1983) copyright Elsevier Publishers B.V.		
	A273	NEBER and RÖCKER, "On the action of benzaldehydes on the free o-aminophenylacetic acid (II)," <u>Chem. Ber.</u> 56:1710-1716 (1923) (GERMAN AND ENGLISH TRANSLATION)		

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Sheet	23	of	31	Prior Application Number	09/741,842
				Prior Appl. Filing Date	12/22/2000
				First Named Inventor	Ken Lipson
				Group Art Unit	Unassigned
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				Attorney Docket Number	038602-1607

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	A277	OSBORNE et al., "Effect of Estrogens and Antiestrogens on Growth of Human Breast Cancer Cells in Athymic Nude Mice," <u>Cancer Research</u> 45:584-590 (1985)			
	A278	O'SULLIVAN and ROTHERY, "The Preparation and Possible Clinical Significance of 4'-Dialkylaminoisoindogenides," <u>Clinica Chimica Acta</u> 62:181-182 (1975) ©Elsevier Scientific Publishing Company			
	A279	OZZELLO and SORDAT, "Behavior of Tumors Produced by Transplantation of Human Mammary Cell Lines in Athymic Nude Mice," <u>Eur. J. Cancer</u> 16:553-559 (1980)			
	A280	PAVLENKO et al., "Introduction of aminomethyl groups into heterocyclic CH-acid molecules," <u>Dopov. Akad. Nauk Ukr Rsr, Ser. B: Geol., Khim. Biol. Nauki</u> 7:64-66 (1980) We should add that we are Sub. Abstract			
	A281	PERKIN et al., "Harmine and Harmaline. Part II. The Synthesis of isoHarman," <u>J. Chem. Soc.</u> 103:1973-1985 (1913)			
	A282	PLATE et al., "Vascular endothelial growth factor is potential tumor angiogenesis factor in human gliomas <i>in vivo</i> ," <u>Nature</u> 359:845-848 (1992)			

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	A284	QUALLICH et al., A General Oxindole Synthesis," <u>J. Synthetic Organic Chemistry</u> : 51-51 (1993)		
	A285	QUINN et al., "Fetal liver kinase 1 is a receptor for vascular endothelial growth factor and is selectively expressed in vascular endothelium," <u>Proc. Natl. Acad. Sci. USA</u> 90:7533-7537 (1993)		
	A286	ROZAKIS-ADCOCK et al., "Association of the Shc and Grb2/Sem5 SH2-containing proteins is implicated in activation of the Ras pathway by tyrosine kinases," <u>Nature</u> 360:689-692 (1992)		
	A287	RUVEDA and GONZALEZ, "Geometric isomerism in benzylideneoxindoles," <u>Spectrochimica Acta</u> 26A:1275-1277 (1970)		
	A288	RYGAARD and POVLSEN, "Heterotransplantation of a Human Malignant Tumour to 'Nude' Mice," <u>Acta Path. Microbiol. Scand.</u> 77:758-760 (1969)		
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	A290	SAITO and STREULI, "Molecular Characterization of Protein Tyrosine Phosphatases," <u>Cell Growth & Differentiation</u> 2:59-65 (1991) ©Molecular Biology Journal of the American Association for Cancer Research		
	A291	SANDBERG-NORDQVIST et al., "Characterization of Insulin-Like Growth Factor 1 in Human Primary Brain Tumors," <u>Cancer Research</u> 53:2475-2478 (1993)		

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	A292	SCHINDLER et al., "Über Dibenz[b,f]-azocin-Derivate," <u>Helvetica Chimica Acta</u> 49:985-989 (1966)			
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	A294	SCHUCHTER et al., "Successful Treatment of Murine Melanoma with Bryostatin 1," <u>Cancer Research</u> 51:682-687 (1991)			
	A295	SEIBERT et al., "Clonal Variation of MCF-7 Breast Cancer Cells <u>in Vitro</u> and in Athymic Nude Mice," <u>Cancer Research</u> 43:2223-2234 (1983)			
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	A298	SHIRAISHI et al., "Specific inhibitors of Tyrosine-Specific Protein Kinase, Synthetic 4-Hydroxycinnamamide Derivatives," <u>Biochemical and Biophysical Research Communications</u> 147:322-328 (1987) © Academic Press			
	A299	SHIRAISHI et al., "Specific Inhibitors of Tyrosine-specific Protein Kinases: Properties of 4-Hydroxycinnamamide Derivatives <u>in Vitro</u> ," <u>Cancer Research</u> 49:2374-2378 (1989)			
	A300	SHWEIKI et al., "Vascular endothelial growth factor induced by hypoxia may mediate hypoxia-initiated angiogenesis," <u>Nature</u> 359:843-845 (1992)			

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	A301	SINGH et al., "Indolinone Derivatives as Potential Antimicrobial Agents," <u>Zentralbl. Mikrobiol.</u> 144:105-109 (1989) copyright VEB Gustav Fischer Verlag Jena		
	A302	SINGH et al., "Synthesis and Anticonvulsant Activity of New 1-Substituted 1'-Methyl-3-Chloro-2-Oxospiro (Azetidin-3', 4-Indol-2' Ones)," <u>Bollettino Chimico Farmaceutico</u> 133:76-79 (1994)		
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	A304	SLAMON et al., "Studies of the HER-2/neu Proto-oncogene in Human Breast and Ovarian Cancer," <u>Science</u> 244:707-712 (1989)		
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	A306	SONGYANG et al., "SH2 Domains Recognize Specific Phosphopeptide Sequences," <u>Cell</u> 72:767-778 (1993) © Cell Press		
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	A308	SPADA, et al., "Small molecule inhibitors of tyrosine kinase activity," <u>Expert Opinion on Therapeutic Patents</u> 5:805-817 (1995) ©Ashley Publications		
	A309	STETINOVA et al., "Stereochemistry and Photoisomerisation of Furfurylideneoxindoles," <u>Collection Czechoslov. Chem. Commun.</u> 42:2201-2206 (1977)		

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	A310	STOLLE, Beilstein Reg. No. 273650, <u>J. Prakt. Chem.</u> , Vol. 2, page 128 (1930)	
	A311	STOLLE, Beilstein Reg. No. 305045, <u>J. Prakt. Chem.</u> , Vol. 2, page 128 (1930)	
	A312	SUMPTER and MILLER, "Chapter IV – Oxindole," in <u>Heterocyclic Compounds With Indole and Carbazole Systems</u> , © Interscience Publishers, Inc., New York, pp. 134-153 (1954)	
	A313	SUN et al., "Design, Synthesis, and Evaluations of Substituted 3-[(3- or 4-Carboxyethylpyrrol-2-yl)methylideny]indolin-2-ones as Inhibitors of VEGF, FGF, and PDGF Receptor Tyrosine Kinases," <u>Journal of Medicinal Chemistry</u> 42: 5120-5130 (1999) ©American Chemical Society	
	A314	SUN et al., "Synthesis and Biological Evaluations of 3-Substituted Indolin-2-ones: A Novel Class of Tyrosine Kinase Inhibitors That Exhibit Selectivity toward Particular Receptor Tyrosine Kinases," <u>J. Med. Chem.</u> 41:2588-2603 (1998) ©The American Chemical Society	
	A315	SUPERTI-FURGA et al., "A functional screen in yeast for regulators and antagonizers of heterologous protein tyrosine kinases," <u>Nature Biotech.</u> 14:600-605 (1996)	
	A316	SUPERTI-FURGA et al., "Csk inhibition of c-Src activity requires both the SH2 and SH3 domains of Src," <u>EMBO J.</u> 12:2625-2634 (1993) © Oxford University Press	
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	A318	TACCONI et al., "(Z)- and (E)-3-Alkylidene-1,3-dihydroindol-2-ones: Influence of Configuration on the Transmission of the Inductive Effect to the Carbonyl Group," <u>J.C.S. Perkin II</u> 150-154 (1976)	

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	A319	TAKANO et al., "Inhibition of angiogenesis by a novel diaminoanthraquinone that inhibits Protein Kinase C," <u>Mol. Bio. Cell</u> 4:358A at abstract no. 2076 (1993)			
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	A326	TUZI et al., "Expression of growth factor receptors in human brain tumours," <u>Br. J. Cancer</u> 63:227-233 (1991)			
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Sheet	30	of	31	Prior Application Number	09/741,842
				Prior Appl. Filing Date	12/22/2000
				First Named Inventor	Ken Lipson
				Group Art Unit	Unassigned
				Examiner Name	Unassigned
				Attorney Docket Number	038602-1607

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(use as many sheets as necessary)				First Named Inv ntor	Ken Lipson
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